

WHAT IS CLAIMED IS:

1. A method for treating or preventing the development of Type II diabetes mellitus in mammals afflicted with such condition with a therapeutically effective amount of a compound of the formula I:

$$R^5$$
 X
 $CH_2OSO_2NHR^1$
 R^2
 R^4
 R^3
 (I)

wherein

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X is CH2 or oxygen;

R1 is hydrogen or alkyl; and

10 R², R³, R⁴ and R⁵ are independently hydrogen or lower alkyl and, when X is CH₂, R⁴ and R⁵ may be alkene groups joined to form a benzene ring and, when X is oxygen, R² and R³ and/or R⁴ and R⁵ together may be a methylenedioxy group of the following formula (II):

$$\begin{array}{c|c}
R^6 & O \\
\hline
R^7 & O
\end{array}$$
(II)

15 wherein

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R⁶ and R⁷ are the same or different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 2. The method of claim 1 wherein the compound of formula (I) is topiramate.
- 3. The method of claim 1, wherein the therapeutically effective amount is from about 10 to 650 mg.
- 4. The method of claim 1, wherein the amount is of from about 16 to 325 mg once or twice daily.
 - 5. A method for treating or preventing the development of Syndrome X (Insulin Resistance Syndrome, Metabolic Syndrome, or Metabolic Syndrome X) in mammals afflicted with such condition with a therapeutically effective amount of a compound of the formula I:





$$R^5$$
 R^4
 R^3
 $CH_2OSO_2NHR^1$
 R^2
 R^3
 (I)

wherein

5

X is CH2 or oxygen;

R1 is hydrogen or alkyl; and

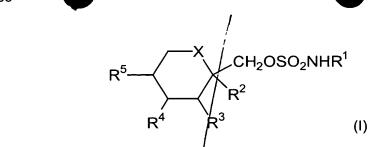
R², R³, R⁴ and R⁵ are independently hydrogen or lower alkyl and, when X is CH₂, R⁴ and R⁵ may be alkene groups joined to form a benzene ring and, when X is oxygen, R² and R³ and/or R⁴ and R⁵ together may be a methylenedioxy group of the following formula (II):

10 wherein

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R⁶ and R⁷ are the same or different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 15 6. The method of claim 5 wherein the compound of formula (I) is topiramate.
 - 7. The method of claim 5, wherein the therapeutically effective amount is from about 10 to 1000 mg daily.
- 20 8. The method of claim 5, wherein the therapeutically effective amount is from about 10 to 650 mg daily.
 - 9. The method of claim 5, wherein the amount is of from about 16 to 325 mg once or twice daily.
- 10. A method for treating impaired oral glucose tolerance in mammals afflicted with such condition with a therapeutically effective amount of a compound of the formula I:



wherein

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X is CH2 or oxygen;

R1 is hydrogen or alkyl; and

R², R³, R⁴ and R⁵ are independently hydrogen or lower alkyl and, when X is CH₂, R⁴ and R⁵ may be alkene groups joined to form a benzene ring and, when X is oxygen, R² and R³ and/or R⁴ and R⁵ together may be a methylenedioxy group of the following formula (II):

10 wherein

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R⁶ and R⁷ are the same of different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 15 11. The method of Claim 10, wherein the compound of formula (I) is topiramate.
 - 12. The method of claim 10, wherein the therapeutically effective amount is from about 10 to 1000 mg daily.
- 20 13. The method of claim 10, wherein the therapeutically effective amount is from about 10 to 650 mg daily.
 - 14. The method of claim 10, wherein the amount is of from about 16 to 325 mg once or twice daily.
- 15. A method for treating or preventing the development of skin lesions associated with Type II diabetes mellitus or Syndrome X in mammals afflicted with such condition with a therapeutically effective amount of a compound of the formula I:



$$R^5$$
 R^4
 R^3
 $CH_2OSO_2NHR^1$
 R^2
 (I)

wherein

5

X is CH2 or oxygen;

R1 is hydrogen or alkyl; and

R², R³, R⁴ and R⁵ are independently hydrogen or lower alkyl and, when X is CH₂, R⁴ and R⁵ may be alkene groups joined to form a benzene ring and, when X is oxygen, R² and R³ and/or R⁴ and R⁵ together may be a methylenedioxy group of the following formula (II):

10 wherein

25

R⁶ and R⁷ are the same or different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 15 16. The method of Claim 15, where in the compound of formula (I) is topiramate.
 - 17. The method of claim 15, wherein the therapeutically effective amount is from about 10 to 1000 mg daily.
- 20 18. The method of claim 15, wherein the therapeutically effective amount is from about 10 to 650 mg daily.
 - 19. The method of claim 15, wherein the amount is of from about 16 to 325 mg once or twice daily.
 - 20. A method for improving defective insulin sensitivity in mammals afflicted with such condition with a therapeutically effective amount of a compound of the formula I:

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$$R^5$$
 R^4
 R^3
 $CH_2OSO_2NHR^1$
 R^2
 (I)

wherein

X is CH2 or oxygen;

R1 is hydrogen or alkyl; and/

R², R³, R⁴ and R⁵ are independently hydrogen or lower alkyl and, when X is CH₂, R⁴ and R⁵ may be alkene groups joined to form a benzene ring and, when X is oxygen, R² and R³ and/or R⁴ and R⁵ together may be a methylenedioxy group of the following formula (II):

$$\begin{array}{c|c}
R^6 & O \\
\hline
R^7 & O \\
\end{array}$$
(II)

10 wherein

R⁶ and R⁷ are the same or different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 21. The method of Claim 20, wherein the compound of formula (I) is topiramate.
 - 22. The method of Claim 20, wherein the therapeutically effective amount is from about 10 to 1000 mg daily.
- 23. The method of claim 20, wherein the therapeutically effective amount is from about 10 to 650 mg daily.
 - 24. The method of claim 20, wherein the amount is of from about 16 to 325 mg once or twice daily.

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